Oncology Clinical Pathways Esophageal Cancer

September 2022 - V1.2022







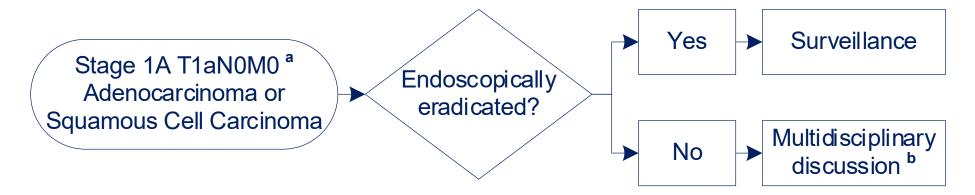
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<u>Esophageal Cancer – Stage 1A T1aN0M0</u> <u>Adenocarcinoma or Squamous Cell Carcinoma</u>



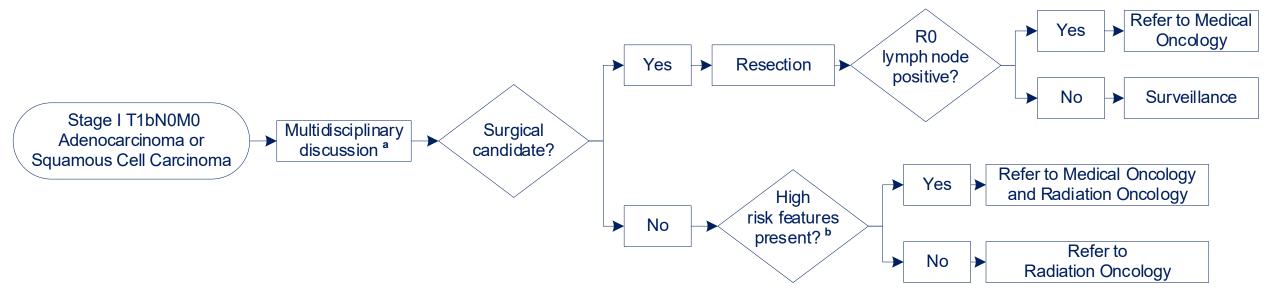
- ^a **Staging** confirmed during Endoscopic Mucosal Resection (EMR) or Endoscopic Submucosal Dissection (ESD)
- ^b Multidisciplinary discussion includes surgery, radiation oncology, and medical oncology







<u>Esophageal Cancer – Stage I T1bN0M0</u> <u>Adenocarcinoma or Squamous Cell Carcinoma</u>



Clinical trial(s) always considered on pathway.

- ^a Multidisciplinary discussion confirm stage
- ^b High risk features include lymphovascular invasion, poorly differentiated tumors, tumor size ≥ 2cm, SM1, SM2, or positive margin on EMR

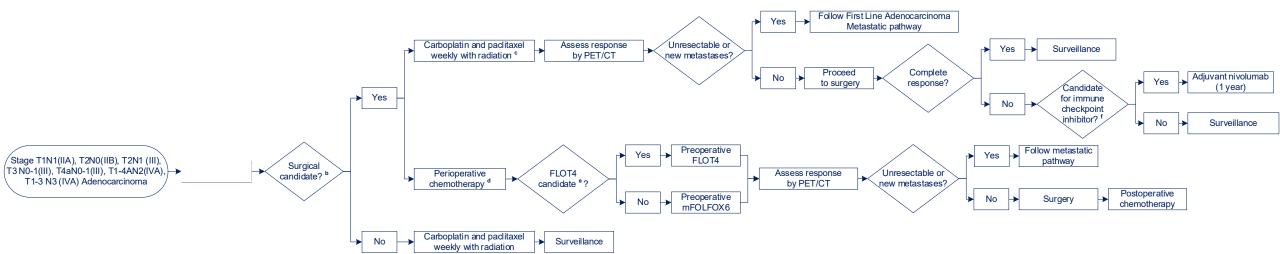
EMR Endoscopic Mucosal Resection







Esophageal Cancer – Stage T1N1(IIA), T2N0(IIB), T2N1 (III), T3 N0-1(III), T4aN0-1(III), T1-4AN2(IVA), T1-3 N3 (IVA) Adenocarcinoma



Clinical trial(s) always considered on pathway

^a Multidisciplinary discussion includes surgery, radiation oncology, and medical oncology

Surgical candidate discussion with or evaluation by a surgeon

Carboplatin and paclitaxel with radiation preferred treatment for predominantly esophageal tumors

Perioperative chemotherapy preferred treatment for predominantly gastric tumors or patient is not a candidate for radiation

e FLOT4 candidates include fit patients with ECOG 0-1 due to expected Grade 3/4 toxicities of neutropenia, infection, diarrhea and peripheral neuropathy

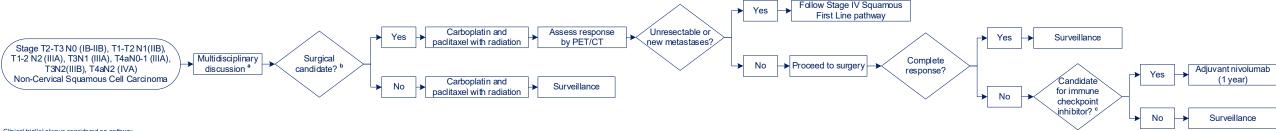
f Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent > 10mg/day) or prior allogeneic HSCT/solid organ transplant







Esophageal Cancer – Stage T2-T3 N0 (IB-IIB), T1-T2 N1(IIB), T1-2 N2 (IIIA), T3N1 (IIIA), T4aN0-1 (IIIA), T3N2(IIIB), T4aN2 (IVA) **Non-Cervical Squamous Cell Carcinoma**



Clinical trial(s) always considered on pathway

^a Multidisciplinary discussion includes surgery, radiation oncology, and medical oncology

Surgical candidate discussion with or evaluation by a surgeon

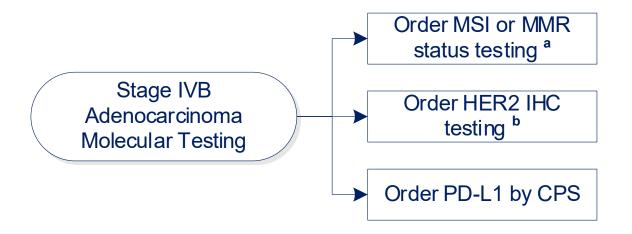
Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant







<u>Esophageal Cancer – Stage IVB Adenocarcinoma Molecular Testing</u>



Clinical trial(s) always considered on pathway.

^a MSI or MMR NGS or IHC

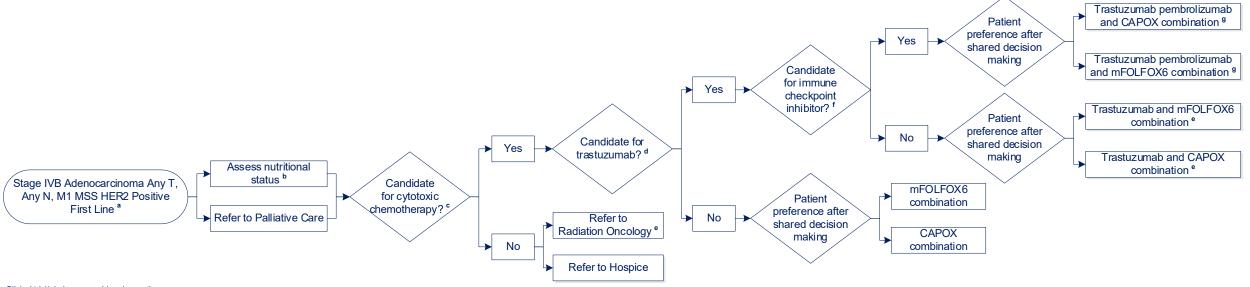
^b **HER2 IHC** if results 2+, HER2 IHC with reflex to FISH required (in-situ hybridization)







<u>Esophageal Cancer – Stage IVB Adenocarcinoma Any T, Any N, M1 MSS HER2 Positive First Line</u>



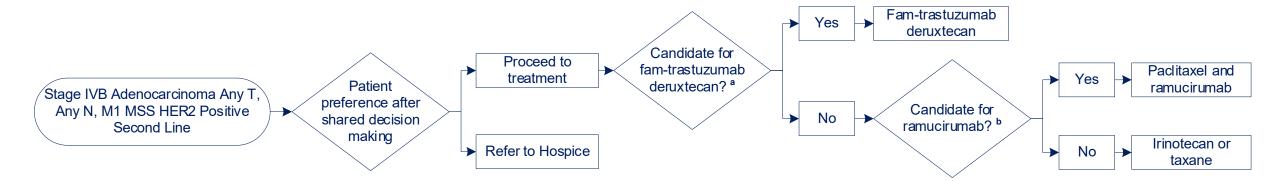
- ^a HER2 Positive considered IHC score +3 or an IHC score of +2 and FISH/ISH positive
- Assess nutritional status and consider palliative stent or other nutritional support modalities when clinically appropriate
- Candidate for cytotoxic chemotherapy consider if patient can tolerate a platinum- and fluoropyrimidine-based doublet
- d Qualify for trastuzumab or biosimilar patient with HER2-positive disease and no clinically significant CV disease (defined as LVEF< 50%, MI within prior 6 months, symptomatic CHF (NYHA class II to IV), unstable angina or cardiac arrhythmia requiring therapy)
- Radiation Oncology consider palliative radiation when clinically appropriate
- Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant
- Pembrolizumab for two years







<u>Esophageal Cancer – Stage IVB Adenocarcinoma Any T, Any N, M1 MSS HER2 Positive Second Line</u>





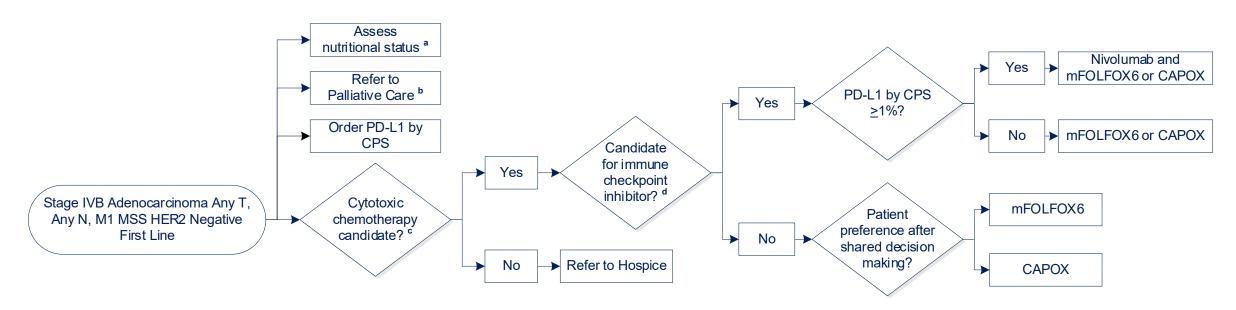




^a Candidate for fam-trastuzumab deruxtecan received trastuzumab in the first-line setting; baseline LVEF ≥ 50% and/or no clinically significant cardiac disease (defined as LVEF < 50%, MI within prior 6 months, symptomatic CHF (NYHA class II to IV), unstable angina or cardiac arrhythmia requiring therapy); no ILD or pneumonitis; ANC ≥ 1500/mm³

b Candidate for ramucirumab received fluoropyrimidine and platinum agent in the first-line setting; ECOG PS 0-2; ANC ≥ 1500/mm³. Note: Due to anti-VEGF effects patients with the following should <u>not</u> receive ramucirumab: non-healing wound/fracture, major surgery in prior 4 weeks, bleeding disorder or coagulopathy, recent history of GI perforation, unstable cardiac condition (uncontrolled HTN, arterial thromboembolism, symptomatic CHF (NYHA II-IV) or arrhythmia), or active cocaine use

<u>Esophageal Cancer – Stage IVB Adenocarcinoma Any T, Any N, M1 MSS HER2 Negative First Line</u>



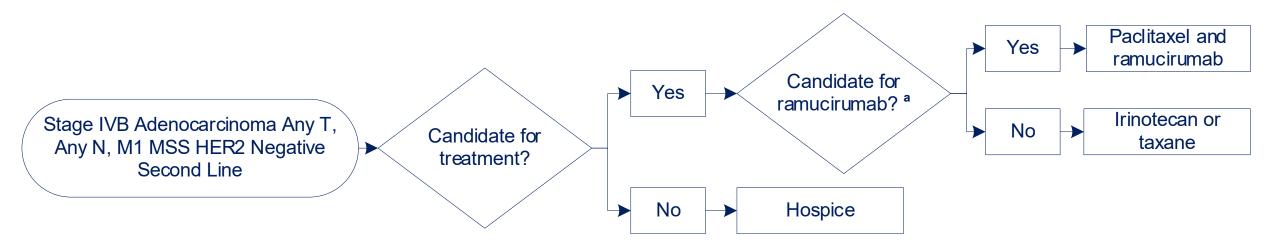
- Assess nutritional status consider palliative stent or other nutritional support modalities when clinically appropriate
- ^b Assess Palliative Care consider palliative radiation when clinically appropriate
- ^c Candidate for cytotoxic chemotherapy consider if patient can tolerate a platinum- and fluoropyrimidine-based doublet
- ^d Qualify for immune checkpoint inhibitor no active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant







<u>Esophageal Cancer – Stage IVB Adenocarcinoma Any T, Any N, M1 MSS HER2 Negative Second Line</u>



Clinical trial(s) always considered on pathway.

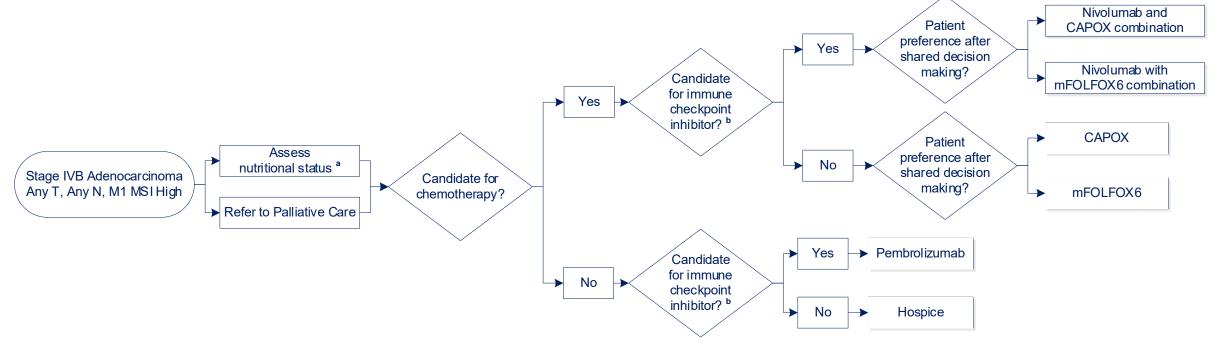
^a Candidate for ramucirumab received fluoropyrimidine and platinum agent in the first-line setting; ECOG PS 0-2; ANC ≥ 1500/mm³; due to anti-VEGF effects patients with the following should <u>not</u> receive ramucirumab: non-healing wound/fracture, major surgery in prior 4 weeks, bleeding disorder or coagulopathy, recent history of GI perforation, unstable cardiac condition (uncontrolled HTN, arterial thromboembolism, symptomatic CHF (NYHA II-IV) or arrhythmia), or active cocaine use







Esophageal Cancer – Stage IVB Adenocarcinoma Any T, Any N, M1 MSI High





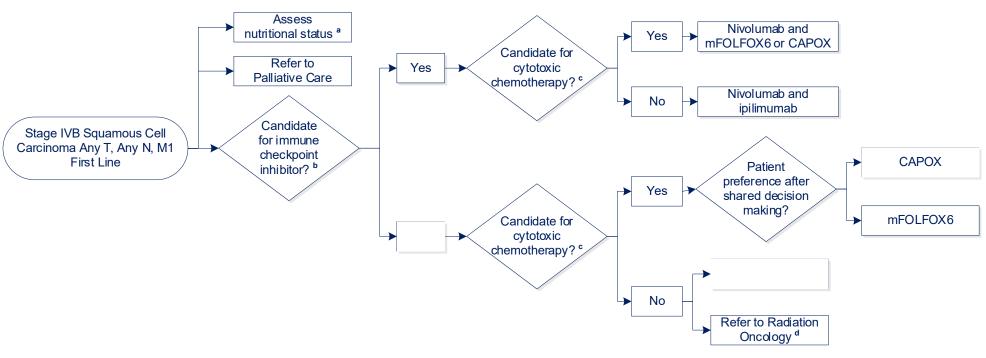




^a Assess nutritional status consider palliative stent or other nutritional support modalities when clinically appropriate

^b Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant

<u>Esophageal Cancer – Stage IVB Squamous Cell Carcinoma</u> <u>Any T, Any N, M1 First Line</u>



d Radiation Oncology consider palliative or metastasis-directed radiation when clinically appropriate





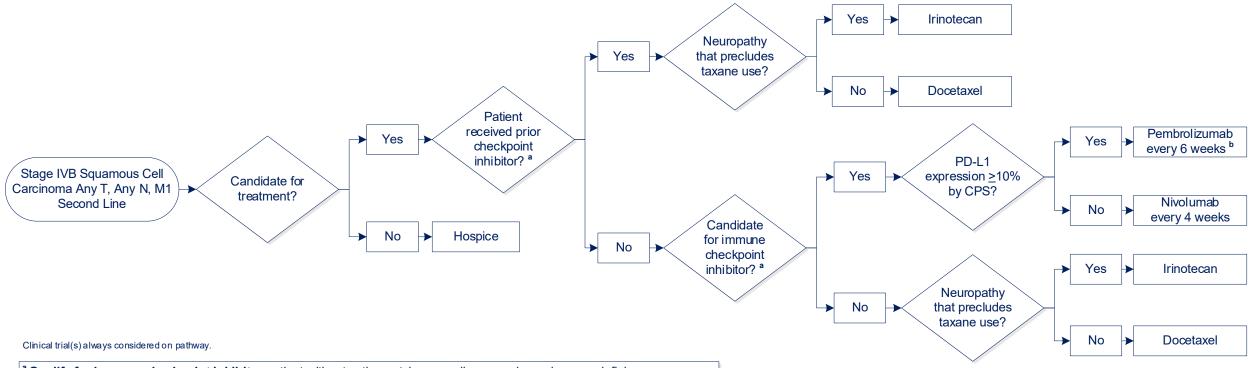


^a Assess nutritional status consider palliative stent or other nutritional support modalities when clinically appropriate

^b Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant

^c Candidate for cytotoxic chemotherapy consider if patient can tolerate a platinum- and fluoropyrimidine-based doublet

<u>Esophageal Cancer – Stage IVB Squamous Cell Carcinoma</u> Any T, Any N, M1 Second Line



^a Qualify for immune checkpoint inhibitor patient without active autoimmune disease, primary immune deficiency, concurrent immunosuppression (including prednisone equivalent >10mg/day) or prior allogeneic HSCT/solid organ transplant

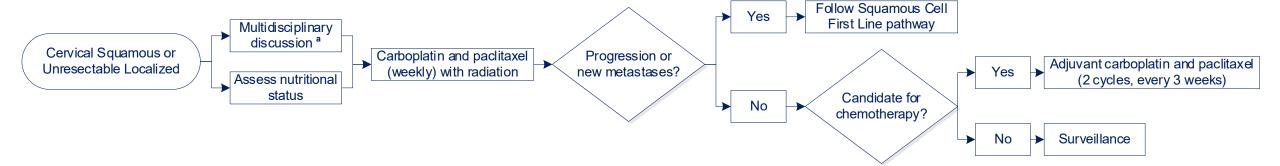
^b Pembrolizumab chosen due to FDA approval and patient convenience







<u>Esophageal Cancer – Cervical Squamous or Unresectable Localized</u>



Clinical trial(s) always considered on pathway.

^a Multidisciplinary discussion includes surgery, radiation oncology, and medical oncology; for mid-esophageal tumors, consider referral to Pulmonary to assess tracheal invasion







Questions?

Contact VHAOncologyPathways@va.gov





